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MICU2 Knockdown 293T cell line, Heterozygous

Catalog No.: RM50185

Basic Information

Catalog No.

RM50185

Category

Cell Lysate

Parental Cell line

293T

Genotype

Knockdown

Background

Enables protein heterodimerization activity. Involved in calcium import into the mitochondrion and negative regulation of mitochondrial calcium ion concentration. Located in mitochondrial inner membrane and mitochondrial intermembrane space. Part of uniplex complex.

Gene Information

Gene Symbol

MICU2

Species

Human

Gene ID

221154

Swiss Prot

Q8IYU8

Synonyms

EFHA1; 1110008L20Rik; MICU2

Contact

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Product Information

Description

MICU2 Knockdown cell line is engineered from 293T cell line with Gene-Editing Technology. Allele-1:105bp deletion in exon1

Allele-2:110bp deletion in exon1

Mammalian cells such as human, rat and mouse cells are normally diploid with two alleles. Homozygote: both alleles were knocked out, mRNA has no signal, no expression of proteins. Heterozygote: only one allele was knocked out, the mRNA transcript levels was decreased compared to wild type, and the protein expression levels was also lower than that of the wild type.

Packaging

1 vial parental cell line and 1 vial knockout cell line

Shipping Conditions

Amount

Dry ice

1~5x10⁶ cells/vial.

Storage

Stored in liquid nitrogen for a long time less than -130°C. Minimizing freeze-thaw cycles.

Protocol

Upon arrival, it should be maintained in DMEM medium with 10%(v/v) fetal bovine serum and 100U penicillin-streptomycin, at 37° C with 5% CO₂ condition.

- 1. Thaw the vial in 37° C water bath ,and shake it to melt as soon as possible.
- 2. Transfer the cell suspension to a 15mL conical tube with pre-warmed 5mL complete medium and centrifuge 1000rpm for approximately 5 minutes at room temperature.
- 3. Remove and discard the supernatant.
- 4. Resuspend the cell pellet with 1mL pre-warmed complete medium and seed in 10cm dish.
- 5. Add 8-10mL of complete medium.
- 6. Incubate the culture at 37°C incubator with 5% CO₂.
- 7. A subcultivation ratio of 1:2-1:4 is recommended.

Sequencing data

WT GGCGGAAAACTGCG***************CGCGTCAGTGTTG
Mut GGCGGAAAACTGCG***Deletion***CCGCGTCAGTGTTG
Allele-1: 105bp deletion in exon1

WT GGCGGAAAACTGCG***************TCAGTGTTGCGGCG
Mut GGCGGAAAACTGCG***Deletion***TCAGTGTTGCGGCG
Allele-2: 110bp deletion in exon1

Genome sequence analysis of PCR products from parental (WT) and MICU2 knockdown (KD) 293T cells, using sanger sequencing.